AMENDMENTS TO THE CLAIMS:

This listing of claims replaces all prior versions and listings of claims in the application:

1-14. (Canceled).

15. (Currently Amended) A method of characterising a treatment applied to a population of cells, comprising:

providing a population of cells;

applying a treatment to the population of cells;

deriving a plurality of cellular features from at least a first captured image of the population of <u>treated</u> cells-that have been exposed to the treatment;

creating an on-target effect signature, which is characteristic of an on-target effect of the treatment on the population of <u>treated</u> cells, from at least a first one of the plurality of cellular features, the at least one of the plurality of features relating to cellular properties involved in the on-target effect;

creating a side effect signature, which is characteristic of a side effect to the ontarget effect of the treatment on the population of treated cells, from at least a second one of the plurality of cellular features, the second one of the plurality of cellular features relating to cellular properties not being involved in the on-target effect;

creating an on-target effect metric derived from the on-target effect signature;

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creating a side effect metric derived from the side effect signature; [[and]]

comparing the on-target effect metric to the side effect metric to derive a ratio of

on-target effect metric to side effect metric to thereby characterize the treatment;

and providing the ratio for a user.

- 16. (Original) The method as claimed in claim 15, wherein the on-target effect signature is created from a group of cellular features.
- 17. (Original) The method as claimed in claim 16, wherein the side effect signature is created from a further group of cellular features, in which none of the members of the group of cellular features used to create the on-target effect signature and the members of the further group of cellular features used to created the side effect signature are common.
- 18. (Original) The method as claimed in claim 15, wherein the second one of the plurality of cellular features is affected by the treatment.
- 19. (Original) The method as claimed in claim 18, further comprising: exposing different populations of cells to different doses of the treatment; and deriving the on-target effect metric and the side effect metric for different doses of the treatment.

- 20. (Previously Presented) The method as claimed in claim 15, wherein deriving one or both of the on-target effect metric and the side effect metric includes determining the difference between the on-target effect signature or side effect signature and a control signature from the same cellular features for a control group of cells.
- 21. (Original) The method as claimed in claim 15, and further comprising: capturing at least a first image of a control group of cells; and deriving a plurality of cellular features from the image of the control group of cells;

creating a control on-target signature for the same cellular features for the control group; and

creating a control side effect signature for the same cellular features for the control group.

- 22. (Original) The method of claim 21, further comprising determining a side effect distance in a multivariate space between the side effect signature and the control side effect signature.
- 23. (Original) The method of claim 22, further comprising determining a target effect distance in a multivariate space between the on-target effect signature and the control on-target effect signature.

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- 24. (Previously Presented) The method of claim 23, wherein characterising the treatment is based on the side effect distance.
- 25. (Previously Presented) The method of claim 24, wherein characterising the treatment is also based on the on-target effect distance.
- 26. (Original) The method as claimed in claim 25, further comprising generating a graphical representation of the side effect distance and on-target effect distance.